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U.S. DEPARTMENT OF COMMERCE **APPLICATION NO.:** ATTY, DOCKET NO .: OC01625K 10/665,005 PATENT AND TRADEMARK OFFICE APPLICANT: INFORMATION DISCLOSURE STATEMENT Kamil Paruch et al. BY APPLICANT GROUP: FILING DATE: (Use several sheets if necessary) 1624 09/19/2003 U.S. PATENT DOCUMENTS FILING DATE IF CLASS DATE NAME SUB-EXAMINER DOCUMENT CLASS APPROPRIATE INITIAL NUMBER AA AB AC AD ΑE AF AG AH ΑI ΑJ AK FOREIGN PATENT DOCUMENTS TRANSLATION SUB-CLASS DATE COUNTRY DOCUMENT CLASS YES NO NUMBER EPO EP 0 778 277 06/11/1997 AL WO 02/06286 01/24/2002 PCT АМ PCT 06/16/1988 AN WO 88/04298 AO AP OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.) Vesely et al., "Inhibition of Cyclin-Dependent Kinases by Purine Analogues", Eur. J. Biochem Senderowicz et al., "Phase I Trial of Continuous Infusion Flavopiridol, a Novel Cyclin-Dependent Kinase Inhibitor, in Patients With Refractory Neoplasms", Journal of Clinical Oncology (September 1998), 16(9): 2986-2999. Meijer et al., "Biochemical and Cellular Effects of Roscovitine, a Potent and Selective Inhibitor of the Cyclin-Dependent Kinases CDC2, CDK2 and CDK5", Eur. J. Biochem. (1997), 243: 527-536. Bible et al., "Cytotoxic Synergy between Flavopiridol (NSC 649890, L86-8275) and Various Antineoplastic Agents: The Importance of Sequence of Administration", Cancer Research (August 15, 1997), **57**: 3375-3380. Shiota et al., "Synthesis and Structure- Activity Relationship of a New Series of Potent Angiotensin II Receptor Antagonists: Pyrazolo[1,5\alpha]pyrimidine Derivatives", Chem. Pharm. Bull. (1999), 47(7): 928-938. Yasuo Makisumi, "Studies on the Azaindolizine Compounds. XI. Synthesis of 6,7-Disubstituted Pyrazolo[1,5-α]pyrimidines.", Chem. Pharm. Bull. (1962), 10: 620-626. DATE CONSIDERED EXAMINER 31004

\*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

- 4